Amendment dated June 23, 2005

In Response to Office Action dated February 23, 2005

## AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in this application.

## Listing of Claims:

Claim 1 (currently amended): A device for inducing local bone or cartilage formation, comprising:

a bone morphogenetic protein selected from the group consisting of OP1, OP2, OP3, BMP2, BMP3, BMP4, BMP5, BMP6, BMP9, BMP10, BMP11, BMP12, BMP15, BMP16, DPP, Vgl, 60A protein, GDF-1, GDF3, GDF5, GDF6, GDF7, GDF8, GDF9, GDF10 and GDF11, capable of inducing repair of endochondral bone, or cartilage, chondral, or osteochondral defects;

a non synthetic, non polymeric matrix selected from the group consisting of collagen, apatites, hydroxyapatites, tricalcium phosphate, and admixtures thereof; and

a binding agent selected from the group consisting of cellulose and salts thereof;

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wherein said binding agent has a degree of substitution of 0.65-0.90 and a viscosity of about 10-200 cP at a 4% (w/v) concentration of said binding agent.

Claim 2 (canceled).

Claim (previously presented): The device of claim 1, wherein said bone morphogenetic protein is selected from the group consisting of OP1, OP2, BMP2, BMP4, BMP5, and BMP6.

Claim (currently amended): A device for inducing local bone and cartilage formation comprising a bone morphogenetic protein comprising an amino acid sequence having at least 70% homology with the C-terminal 102-106 amino acids, including the conserved seven cysteine domain, of human OP1;

a non-synthetic, non-polymeric matrix selected from the group consisting of collagen, apatites, hydroxyapatites, tricalcium phosphate, and admixtures thereof; and

a binding agent selected from the group consisting of cellulose and salts thereof;

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wherein said binding agent has a degree of substitution of 0.65-0.90 and a viscosity of about 10-200 cP at a 4% (w/v) concentration of said binding agent; and

wherein said bone morphogenetic protein is capable of inducing repair of endochondral bone when implanted together with a matrix in a mammal.

Claim & (previously presented): The device of claim 1 wherein said bone morphogenetic protein is OP-1.

Claim (withdrawn): The device of claim 1 wherein said device comprises at least two different bone morphogenetic proteins.

Claim 7 (canceled).

Claim (original): The device of claim 1 wherein said matrix is collagen.

Claim (withdrawn): The device of claim 1 wherein said device comprises at least two different matrix materials.

Claim 10 (canceled).

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Claim 1 (currently amended): The device of claim 1 wherein said binding agent is selected from the group consisting of alkylcelluloses an alkylcellulose.

Claim 12 (original): The device of claim 1 wherein said binding agent is selected from the group consisting of methylcellulose, methylhydroxyethylcellulose, hydroxyethylcellulose, hydroxypropylmethylcellulose, carboxymethylcellulose, sodium carboxymethylcellulose, hydroxyalkylcelluloses, and admixtures thereof.

claim 12 (currently amended): The device of claim 1 wherein said binding agent is carboxymethylcellulose or the a sodium salt thereof.

Claim 1 (withdrawn): The device of claim 1 wherein said device comprises at least two different binding agents.

Claim 18 (original): The device of claim 1 further comprising a wetting agent.

Claim 16 (original): The device of claim 1/5 wherein said wetting agent is saline.

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Claim 1/1 (currently amended): A device for inducing local bone or cartilage formation, comprising at least approximately 1.25 mg of OP-1 and at least approximately 180 mg of carboxymethylcellulose per 1000 mg of collagen matrix, wherein said carboxymethylcellulose has a degree of substitution of 0.65-0.90 and a viscosity of about 10-200 cP at a 4% (w/v) (w/v) concentration of said carboxymethylcellulose.

Claim 18 (previously presented): The device of claim 19 comprising at least approximately 2.5 mg of OP-1 per 1000 mg of collagen matrix.

Claim 18 (previously presented): The device of claim 18 or 18 comprising at least approximately 200 mg of carboxymethylcellulose per 1000 mg of collagen matrix.

Claim 20 (previously presented): The device of claim 1 wherein the binding agent to matrix ratio is one part by weight binding agent to 1-10 parts by weight matrix.

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Claim 21 (previously presented): The device of claim 20 wherein the binding agent to matrix ratio is one part by weight binding agent to 5 parts by weight matrix.

Claim 22 (previously presented): The device of claim 22 wherein the binding agent to matrix ratio is one part by weight binding agent to 1-5 parts by weight matrix.

Claim 23 (previously presented): The device of claim 1 wherein the binding agent to matrix ratio is 1-10 parts by weight binding agent to 1 part by weight matrix.

Claim 2# (previously presented): The device of claim 2/3 wherein the binding agent to matrix ratio is fewer than 10 parts by weight binding agent to one part by weight matrix.

Claim 25 (currently amended): The device of claim  $1/\sqrt{7}$ or 16 or 19 further comprising saline.

Claims 26-30 (canceled).

Claim 31 (previously presented): A device for inducing local bone or cartilage formation comprising:

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OP-1;

collagen matrix; and

carboxymethylcellulose having a degree of substitution of 0.65-0.90 and a viscosity of about 10-200 cP at a 4% (w/v) concentration of said carboxymethylcellulose.

Claim 32 (currently amended): A kit for inducing local bone or cartilage formation, the kit comprising:

- (a) a first receptacle housing a bone morphogenetic protein selected from the group consisting of OP1, OP2, OP3,

  BMP2, BMP3, BMP4, BMP5, BMP6, BMP9, BMP10, BMP11, BMP12, BMP15,

  BMP16, DPP, Vgl, 60A protein, GDF-1, GDF3, GDF5, GDF6, GDF7,

  GDF8, GDF9, GDF10 and GDF11 and a non synthetic, non polymeric matrix selected from the group consisting of collagen, apatites, hydroxyapatites, tricalcium phosphates and admixtures thereof, and
- (b) a second receptacle housing a binding agent selected from the group consisting of cellulose, and salts thereof,

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wherein said binding agent has a degree of substitution of 0.65-0.90 and a viscosity of about 10-200 cP at a 4% (w/v) concentration of said binding agent.

Claim 3/3 (previously presented): The kit of claim 3/2 further comprising a receptacle adapted to house a wetting agent.

Claim 34 (canceled).

Claim 36 (currently amended): A kit for inducing local bone or cartilage formation, the kit comprising:

a first receptacle housing a bone morphogenetic protein selected from the group consisting of OP1, OP2, OP3, BMP2, BMP3, BMP4, BMP5, BMP6, BMP9, BMP10, BMP11, BMP12, BMP15, BMP16, DPP, Vgl, 60A protein, GDF-1, GDF3, GDF5, GDF6, GDF7, GDF8, GDF9, GDF10 and GDF11, a non-synthetic, non-polymeric matrix selected from the group consisting of collagen, apatites, hydroxyapatites, tricalcium phosphates, and admixtures thereof, and a binding agent selected from the group consisting of cellulose, and salts thereof,

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wherein said binding agent has a degree of substitution of 0.65-0.90 and a viscosity of about 10-200 cP at a 4% (w/v) concentration of said binding agent.

Claim 36 (previously presented): The kit of claim 36, further comprising a second receptacle adapted to house a wetting agent.

Claim 3/ (previously presented): The device of claim/5, wherein the amount of the OP-1 ranges from approximately 0.125 mg to 10.0 mg.

Claim 38 (previously presented): The device of claim

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37, wherein the amount of the OP-1 is approximately 3.5 mg.

Claim 35 (new): The device of claim 15 further comprising saline.